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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)			
	10/537,116	GRINVALD ET AL.			
Office Action Summary	Examiner	Art Unit			
	KATHERINE L. FERNANDEZ	3768			
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address			
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).			
Status					
 1) ☐ Responsive to communication(s) filed on <u>05 Fe</u> 2a) ☐ This action is FINAL. 2b) ☐ This 3) ☐ Since this application is in condition for allowar closed in accordance with the practice under E 	action is non-final. nce except for formal matters, pro				
Disposition of Claims					
4) Claim(s) 45-56,58 and 66-81 is/are pending in 4a) Of the above claim(s) is/are withdray 5) Claim(s) is/are allowed. 6) Claim(s) 45-56,58 and 66-81 is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) are subject to restriction and/or Application Papers	vn from consideration.				
9) ☐ The specification is objected to by the Examiner 10) ☑ The drawing(s) filed on 02 June 2005 is/are: a) Applicant may not request that any objection to the of Replacement drawing sheet(s) including the correction 11) ☐ The oath or declaration is objected to by the Examiner	☑ accepted or b)☐ objected to drawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 					
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 1/21/09.	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	ate			

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Claim Rejections - 35 USC § 103

1. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

- (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 2. Claim 45-50, 53, 66-67, 70-77, 79-81 are rejected under 35 U.S.C. 103(a) as being unpatentable over Grinvald et al. (WO 99/63882) as cited by applicant in view of Norris et al. (US Patent No. 5,477,858).

Grinvald et al. disclose a method for vascular analysis of a subject, comprising the steps of: optically imaging moving erythrocytes within at least one optically accessible blood vessel of a subject (pg. 3, last paragraph-pg. 4, top paragraph; pg. 5; pg. 8, 1st-2nd paragraphs); determining from said optical imaging at least one flow characteristic of said erythrocytes in at least one optically accessible blood vessel (pg. 5; pg. 9, 5th paragraph-pg. 10, 1st paragraph; pg. 12, 3rd paragraph, referring to determining blood flow direction and a rate-of-flow map from data from images) and generating an output on an output device (pg. 5). The blood vessel can be a retinal blood vessel or can be located in tissue of an internal organ (i.e. brain tissue) (pg. 7, last paragraph-pg. 8, first paragraph). The optical imaging comprises acquiring at least two sequential images of erythrocytes in said at least one optically accessible blood vessel (pg. 3, last paragraph-pg. 4, 2nd paragraph; pg. 5, last paragraph). Grinvald et al. further disclose a system for performing the method discussed above, which includes a light source for illuminating at least one optically accessible blood vessel of the subject

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(pg. 6, 3rd paragraph), an imager for acquiring a plurality of images of moving erythrocytes showing sequential spatial distribution of said moving erythrocytes in said at least one optically (pg. 5), an image discriminator for determining from said plurality of images showing sequential spatial distribution, a flow pattern of erythrocytes along said blood vessel (pg. 5; pg. 9, last paragraph-pg. 10, 2nd paragraph), and a flow analyzer for analyzing said flow pattern to determine at least one flow characteristic of erythrocytes along said at least one optically accessible blood vessel of the subject (pg. 5; pg. 11, 2nd paragraph; pg. 12, 3rd-4th paragraph). Their system further comprises a wavelength selector (i.e. bandpass filter) configured to configure said imager to acquire said images of said at least one optically accessible blood vessel over a limited wavelength band (pg. 6, last paragraph). The wavelength selector is located in an illuminating pathway between said light source and said at least one optically accessible blood vessel and in an imaging pathway between said at least one optically accessible blood vessel and said imager (pg. 6, last paragraph). Although Grinvald et al. do not specifically disclose that the wavelength band is between 2 and 30 nm, it would have been within the skill of one of ordinary skill in the art to modify the invention of Grinvald et al. to experimentally adjust the wavelength band to between 2 and 30 nm in order to determine the appropriate wavelength. The light source for illuminating said at least one optically accessible blood vessel of the subject is a pulsed source having a pulse to pulse interval of less than 1 second or between 5 and 200 ms (pg. 5, last paragraph-pg. 6, 3rd paragraph). Grinvald et al. further disclose that their system includes a computer

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and a display monitor for viewing the results of automatic image analysis and permitting interactive image analysis, and a printer for hard copy output of analysis results (pg. 5).

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However, they do not specifically disclose that their method includes the step of utilizing said at least one flow characteristic for identifying roughness on an inner wall of said at least one optically accessible blood vessel or that their invention includes a wall analyzer for utilizing said at least one flow characteristic for determining at least one property of an inner surface wall of said blood vessel.

Norris et al. disclose an ultrasound imaging system for creating real-time images of tissue and blood flow (column 1, lines 18-21). They disclose that when plaque build up in vessels (i.e. causing roughness on an inner wall of vessel), the direction and speed of blood flow are altered (column 1, line 62-column 2, line 4). At the time if the invention, it would have been obvious to one of ordinary skill in the art to modify the invention of Grinvald et al. to have their computer serve as a wall analyzer to perform the step of utilizing at least one flow characteristic (i.e. flow direction, flow rate) for identifying roughness on an inner wall of said at least one optically accessible blood vessel, as Norris teaches that flow characteristics (i.e. flow direction, flow rate) are indicators of plaque formation (i.e. roughness on inner wall of blood vessel) (column 1, line 62-column 2, line 4).

3. Claims 51-52 and 68-69 are rejected under 35 U.S.C. 103(a) as being unpatentable over Grinvald et al. in view of Norris et al. as applied to claims 45 and 67 above, and further in view of Wong et al. ("Retinal microvascular abnormalities and

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incident stroke: the Athersclerosis Risk in Communities Study", October 2001) as cited by applicant.

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As discussed above, the combined references of Grinvald et al. in view of Norris et al. meet the limitations of claims 45 and 67. However, they do not specifically disclose that their method includes the step of utilizing said identifying of said roughness on said inner wall of said at least one optically accessible blood vessel in order to determine a level of arteriosclerosis in the subject or that their system. Further, they do not specifically disclose that their method further comprises the step of utilizing said identifying of said roughness on said inner wall of said at least one optically accessible blood vessel in order to determine a condition of another blood vessel of the subject. Wong et al. disclose a study investigating the relation between retinal microvascular abnormalities to incident stroke (pg. 1134, left column, Summary:Background). They conclude that retinal microvascular lesions (i.e. roughness on wall of blood vessel) can be markers of general vascular pathology (such as atherosclerosis, which is a form of arteriosclerosis) rather than specific microvascular pathology (pg. 1139, left column, 1st paragraph). At the time of the invention, it would have been obvious to one of ordinary skill in the art to have included the above limitations to the method of Grinvald et al. in view of Norris et al., as taught by Wong et al., in order to provide a non-invasive way of determining the risk of arteriosclerosis in an individual (pg. 1139, left column, 1st paragraph).

4. Claims 54-56 and 58 are rejected under 35 U.S.C. 103(a) as being unpatentable over Grinvald et al. in view of Norris et al., and further in view of Taylor '02 ("In Vivo

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Quantification of Blood Flow and Wall Shear Stress in the Human Abdominal Aorta During Lower Limb Exercise", March 2002).

As discussed above, the combined references of Grinvald et al. and Norris et al. meet most of the limitations of claim 54 (i.e. optically imaging moving erythrocytes once and then again, determining from said optical imaging at least one erythrocytic flow characteristic, and utilizing differences in said at least one flow characteristic to determine a roughness index of an inner wall of said at least one optically accessible blood vessel). However, they do not specifically disclose that optically imaging the moving erythrocytes the first time is performed with subject having a first blood pressure, said first pressure being subject to change to a second blood pressure, optically imaging the moving erythrocytes within the blood vessel again when said first blood pressure of said subject has changed to said second blood pressure, nor do they disclose that the change from said first blood pressure to said second blood pressure is caused by at least one of exercise and drugs administered to the subject. They also do not disclose that the change of said first blood pressure to said second blood pressure is a result of the subject's heartbeat. Further, they do not disclose that their method includes the limitations of instant claims 56 and 58, which disclose that the first blood pressure corresponds to a first point in a cardiac cycle of the subject wherein said second blood pressure corresponds to a second point in the cardiac cycle of the subject, and wherein the optical imaging steps comprise optically imaging moving erythrocytes within said at least one optically accessible blood vessel when the subject's cardiac cycle is respectively at said first and second points in the subject's cardiac

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cycle, or that the optical imaging steps comprise detecting a parameter of the subject selected from the group consisting of the subject's cardiac cycle and blood pressure of the subject, and optically imaging the moving erythrocytes in response to the selected parameter.

Taylor '02 disclose a study that measures, in vivo, the spatial distribution of blood flow velocities in the abdominal aorta of human subjects during upright rest and light exercise conditions (pg. 403, left column, 3rd paragraph). They disclose that data was collected at rest and during steady-state exercise conditions within the range of light exercises (i.e. data was collected at two different heart rates, which cause a change in blood pressure; also scans were synchronized with the subject's heart beat) (pg. 403, right column, 2nd paragraph). They further disclose that the image acquisitions were gated to the cardiac cycle using a plethysmograph, and that the data was retrospectively reconstructed at 16 discrete time points within the cardiac cycle (pg. 403, right column, 3rd paragraph). Further, the subjects monitored their own heart rate, which was displayed in real-time on a pulse monitor (pg. 403, right column, 2nd paragraph). At the time of the invention, it would have been obvious to one of ordinary skill in the art to have the optical imaging steps in the method of Grinvald et al. in view of Norris et al. be performed at different blood pressure readings, wherein the change in pressure readings is a result of exercise performed by the subject and to further include the limitations of claims 56 and 58, as taught by Taylor '02, in order to determine the effect of activities that change blood pressure (such as exercise) have on flow characteristics (pg. 403, left column, 2nd-3rd paragraphs).

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5. Claim 78 is rejected under 35 U.S.C. 103(a) as being unpatentable over Grinvald et al. in view of Norris et al. as applied to claim 66 above, and further in view of Flower '94 (US Patent No. 5,279,298).

As discussed above, the above combined references meet the limitations of claim 66. Further, they disclose that the imager is configured to acquire images at predetermined intervals (see Grinvald, pg. 5, last paragraph-pg. 6, 2nd paragraph). However, they do not specifically disclose that the light source for illuminating said at least one optically accessible blood vessel of the subject is a continuous source. Flower '94 discloses a method and apparatus to detect and treat neovascular membranes in the ocular vasculature of the fundus of the eye (column 1, lines 8-10). They disclose the use of a continuous light source and that the imager acquires the images at predetermined intervals (column 5, lines 34-52, referring to the computer recording successive images or frames of the fundus of the eye with the passage of time). At the time of the invention, it would have been obvious to one of ordinary skill in the art to include the above limitations in the system of Grinvald et al. in view of Norris et al., as their invention requires an illumination source and Flower et al. teach the successful use of a continuous illumination source when imaging a vessel.

Response to Arguments

6. Applicant's arguments with respect to claims 45-56, 58 and 66-81 have been considered but are most in view of the new ground(s) of rejection.

Conclusion

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7. Any inquiry concerning this communication or earlier communications from the examiner should be directed to KATHERINE L. FERNANDEZ whose telephone number is (571)272-1957. The examiner can normally be reached on 8:30-5, Monday-Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on (571)272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Eric F Winakur/ Primary Examiner, Art Unit 3768